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Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

- 1-5. (Canceled)
- 6. (Currently Amended) A method for treating a gastrointestinal disorder; for treating and/or improving a gastrointestinal property of a COX-2 selective inhibitor; for decreasing the recurrence of an ulcer; for improving a gastroprotective property, an anti-Helicobacter pylori property or an antacid property of a proton pump inhibitor; or for improving a gastroprotective property of an H₂ receptor antagonist; in a patient in need thereof comprising administering to the patient a therapeutically effective amount of:

N-nitrato-pivaloyl-S-(N-acetyl-glycyl)-L-cysteine ethyl ester (compound SPM 5186) or a pharmaceutically acceptable salt thereof; N-nitrato-pivaloyl-S-(N-acetyl-alanyl)-L-cysteine ethyl ester (compound SPM 5185) or a pharmaceutically acceptable salt thereof; N-nitrato-pivaloyl-S-(N-acetyl-leucyl)-L-cysteine ethyl ester; N-(2-nitratoacetyl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoacetyl)-S-acetyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoacetyl)-S-propionyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoacetyl)-S-pivaloyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoacetyl)-methionine methyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratopropionyl)-cysteine or a pharmaceutically acceptable salt thereof; N-(2-nitratopropionyl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratopropionyl)-methionine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratobutyryl)-cysteine or a pharmaceutically acceptable salt thereof; N-(2-nitratobutyryl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratobutyryl)-S-acetyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratobutyryl)-S-butyryl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratobutyryl)-methionine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoisobutyryl)-cysteine or a pharmaceutically acceptable salt thereof; N-(2nitratoisobutyryl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2nitratoisobutyryl)-S-benzoyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoisobutyryl)-S-acetyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratoisobutyryl)-S-pivaloyl-cysteine ethyl ester or a pharmaceutically-acceptable

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salt-thereof; N-(2-nitratoisobutyryl)-methionine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratobutyryl)-cysteine or a pharmaceutically acceptable salt thereof; N-(3nitratobutyryl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3nitratobutyryl)-S-acetyl-cysteine ethyl ester or a pharmaceutically acceptable salt-thereof; N-(3nitratobutyryl)-S-propionyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratobutyryl)-methionine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3nitratobutyryl)-homocysteine thiolactone or a pharmaceutically acceptable salt thereof; N-(3nitratopivaloyl)-cysteine or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)cysteine ethyl ester-S-ethyl carbonate or a pharmaceutically acceptable salt thereof; N-(3nitratopivaloyl)-S-acetyl-cysteine ethyl ester or a pharmaceutically-acceptable salt thereof; N-(3nitratopivaloyl)-S-propionyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-S-butyryl-cysteine ethyl ester or a pharmaceutically acceptable salt-thereof; N-(3-nitratopivaloyl)-S-isobutyryl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-S-pivaloyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-S-benzoyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-methionine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-methionine or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-homocysteine thiolactone or a pharmaceutically acceptable salt thereof; N-(2-nitratohexanoyl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(2-nitratohexanoyl)-S-propionyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratohexanoyl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(3-nitratohexanoyl)-methionine methyl ester or a-pharmaceutically acceptable salt-thereof; N-(12-nitratolauroyl)-cysteine or a pharmaceutically acceptable salt thereof; N-(12nitratolaurovl)-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; N-(12nitratolauroyl)-S-acetyl-cysteine or a pharmaceutically acceptable salt thereof; N-(12nitratolauroyl)-S-pivaloyl-cysteine or a pharmaceutically acceptable salt-thereof; or compound SPM-6373 N-(3-hydroxy-2,2-dimethyl-1-oxopropyl)-3-[(acetylamino)acetate]- L-cysteine ethyl ester (SPM 6373); or a pharmaceutically acceptable salt thereof.

7. (Previously Presented) The method of claim 6, further comprising administering a pharmaceutically acceptable carrier.

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8. (Previously Presented) The method of claim 6, further comprising administering an NSAID, a COX-2 inhibitor, an H₂ receptor antagonist, a proton pump inhibitor, a vasoactive agent, a steroid, a ß-agonist, an anticholinergic, a mast cell stabilizer, a PDE inhibitor, taxane, rapamycin, tranilast, or a combination of two or more thereof.

9. (Cancelled)

- 10. (Currently Amended) The method of claim 6, comprising administering to the patient N-nitrato-pivaloyl-S-(N-acetyl-glycyl)-L-cysteine ethyl ester (compound SPM 5186) or a pharmaceutically acceptable salt thereof; N-nitrato-pivaloyl-S-(N-acetyl-alanyl)-L-cysteine ethyl ester (compound SPM 5185) or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-S-pivaloyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; or compound SPM 6373 N-(3-hydroxy-2,2-dimethyl-1-oxopropyl)-3-[(acetylamino)acetate]- L-cysteine ethyl ester (SPM 6373); or a pharmaceutically acceptable salt thereof.
- 11. (Previously Amended) The method of claim 6, wherein the method is the method for treating a gastrointestinal disorder.
- 12. (Previously Amended) The method of claim 6, wherein the method is the method for treating and/or improving a gastrointestinal property of a COX-2 selective inhibitor.
- 13. (Previously Presented) The method of claim 6, wherein the method is the method for decreasing the recurrence of an ulcer.
- 14. (Previously Presented) The method of claim 6, wherein the method is the method for improving a gastroprotective property of a proton pump inhibitor.
- 15. (Previously Presented) The method of claim 6, wherein the method is the method for improving an anti-*Helicobacter pylori* property of a proton pump inhibitor.
- 16. (Previously Presented) The method of claim 6, wherein the method is the method for improving an antacid property of a proton pump inhibitor.
- 17. (Previously Presented) The method of claim 6, wherein the method is the method for improving a gastroprotective property of an H₂ receptor antagonist.
- 18. (Currently Amended) A method for treating a gastrointestinal disorder; for treating and/or improving a gastrointestinal property of a COX-2 selective inhibitor; for decreasing the recurrence of an ulcer; for improving a gastroprotective property, an anti-*Helicobacter pylori* property or an antacid property of a proton pump inhibitor; or for improving a gastroprotective property of an H₂ receptor antagonist in a patient in need thereof comprising administering to the

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patient a therapeutically effective amount of at least one compound selected from the group consisting of N-nitrato-pivaloyl-S-(N-acetyl-glycyl)-L-cysteine ethyl ester (compound-SPM 5186) or a pharmaceutically acceptable salt thereof; N-nitrato-pivaloyl-S-(N-acetyl-alanyl)-L-cysteine ethyl ester (compound SPM 5185) or a pharmaceutically acceptable salt thereof; N-(3-nitratopivaloyl)-S-pivaloyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof; and compound SPM 6373; N-(3-hydroxy-2,2-dimethyl-1-oxopropyl)-3-[(acetylamino)acetate]- L-cysteine ethyl ester (SPM 6373); or a pharmaceutically acceptable salt thereof.

- 19. (Previously Presented) The method of claim 18, further comprising administering a pharmaceutically acceptable carrier.
- 20. (Previously Presented) The method of claim 18, further comprising administering an NSAID, a COX-2 inhibitor, an H₂ receptor antagonist, a proton pump inhibitor, a vasoactive agent, a steroid, a β-agonist, an anticholinergic, a mast cell stabilizer, a PDE inhibitor, taxane, rapamycin, tranilast, or a combination of two or more thereof.
- 21. (Currently Amended) The method of claim 18, comprising administering to the patient a therapeutically effective amount of N-nitrato-pivaloyl-S-(N-acetyl-glycyl)-L-cysteine ethyl ester (compound SPM 5186) or a pharmaceutically acceptable salt thereof.
- 22. (Currently Amended) The method of claim 18, comprising administering to the patient a therapeutically effective amount of N-nitrato-pivaloyl-S-(N-acetyl-alanyl)-L-cysteine ethyl ester (compound SPM 5185) or a pharmaceutically acceptable salt thereof.
- 23. (Previously Presented) The method of claim 18, comprising administering to the patient a therapeutically effective amount of N-(3-nitratopivaloyl)-S-pivaloyl-cysteine ethyl ester or a pharmaceutically acceptable salt thereof.
- 24. (Currently Amended) A method for treating and/or improving a gastrointestinal property of a COX-2 selective inhibitor; for decreasing the recurrence of an ulcer; for improving a gastroprotective property, an anti-Helicobacter pylori property or an antacid property of a proton pump inhibitor; or for improving a gastroprotective property of an H₂ receptor antagonist in a patient in need thereof comprising administering to the patient a therapeutically effective amount of N-(3-nitratopivaloyl)-1-cysteine ethyl ester (SPM 3672);compound SPM 3672 or a pharmaceutically acceptable salt thereof.
- 25. (Currently Amended) The method of claim 18, comprising administering to the patient a therapeutically effective amount of compound SPM 6373 N-(3-hydroxy-2,2-dimethyl-

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1-oxopropyl)-3-[(acetylamino)acetate]- L-cysteine ethyl ester (SPM 6373); or a pharmaceutically acceptable salt thereof.